

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product datasheet for RC216151L3V

Von Hippel Lindau (VHL) (NM_000551) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type:	Lentiviral Particles	
Product Name:	Von Hippel Lindau (VHL) (NM_000551) Human Tagged ORF Clone Lentiviral Particle	
Symbol:	Von Hippel Lindau	
Synonyms:	HRCA1; pVHL; RCA1; VHL1	
Mammalian Cell Selection:	Puromycin	
Vector:	pLenti-C-Myc-DDK-P2A-Puro (PS100092)	
Tag:	Myc-DDK	
ACCN:	NM_000551	
ORF Size:	639 bp	
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC216151).	
OTI Disclaimer:	The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <u>More info</u>	
OTI Annotation:	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.	
RefSeq:	<u>NM 000551.2</u>	
RefSeq Size:	2968 bp	
RefSeq ORF:	642 bp	
Locus ID:	7428	
UniProt ID:	<u>P40337</u>	
Cytogenetics:	3p25.3	
Domains:	VHL	
Protein Families:	Druggable Genome, Transcription Factors	



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

	Von Hippel Lindau (VHL) (NM_000551) Human Tagged ORF Clone Lentiviral Particle – RC216151L3V	
Protein Pathwa	ys: Path	ways in cancer, Renal cell carcinoma, Ubiquitin mediated proteolysis
MW: 24 kDa		Da
Gene Summary	pred is the com poss degr cent POLI	Hippel-Lindau syndrome (VHL) is a dominantly inherited familial cancer syndrome disposing to a variety of malignant and benign tumors. A germline mutation of this gene e basis of familial inheritance of VHL syndrome. The protein encoded by this gene is a ponent of the protein complex that includes elongin B, elongin C, and cullin-2, and esses ubiquitin ligase E3 activity. This protein is involved in the ubiquitination and adation of hypoxia-inducible-factor (HIF), which is a transcription factor that plays a ral role in the regulation of gene expression by oxygen. RNA polymerase II subunit R2G/RPB7 is also reported to be a target of this protein. Alternatively spliced transcript ants encoding distinct isoforms have been observed. [provided by RefSeq, Jul 2008]

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US